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Process for activating a hydrotreating catalyst

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PROCESS FOR ACTIVATING A HYDROTREATING CATALYST

- The invention pertains to a process for activating a hydrotreating catalyst. The hydrotreating catalyst to be activated may be either a fresh hydrotreating catalyst or a hydrotreating catalyst which has been used and regenerated. The present invention also pertains to the hydrotreating catalyst thus obtained, and to its use in hydrotreating.
- 10 In general, the object of catalytically hydrotreating hydrocarbon-containing feeds is the removal of impurities. Common impurities are sulfur compounds and nitrogen compounds. The at least partial removal of such impurities from a feed will ensure that, when the final product is burnt, fewer sulfur oxides and/or nitrogen oxides harmful to the environment will be released. In addition, sulfur compounds and nitrogen compounds are toxic to many of the catalysts employed in the oil industry for converting feeds into ready-for-use products. Examples of such catalysts include cracking catalysts, hydrocracking catalysts, and reforming catalysts. It is therefore customary for feeds to be subjected to a catalytic hydrotreatment prior to their being processed in, say, a cracking unit.
- 20 Catalytic hydrotreatment implies contacting a feed with hydrogen at elevated temperature and pressure in the presence of a hydrotreating catalyst. In this process the sulfur compounds and nitrogen compounds present in the feed are converted into readily removable hydrogen sulfide and ammonia.
- 25 In general, hydrotreating catalysts are composed of a carrier with deposited thereon a Group VI metal component and a Group VIII metal component. The most commonly employed Group VI metals are molybdenum and tungsten, while cobalt and nickel are the conventional Group VIII metals. Phosphorus may also be present in the catalyst. The prior art processes for preparing these catalysts are characterized in that a carrier material is composited with hydrogenation metal components, for example by impregnation, after which the composite is calcined to convert the metal components into their oxides. Before
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being used in hydrotreating, the catalysts are generally presulfided to convert the hydrogenation metals into their sulfides.

Because the requirements as to the legally permitted sulfur and nitrogen contents in fuels are becoming ever stricter, there is a continuous need for
5 hydrotreating catalysts with improved activity. Further, at a given final sulfur content a more active catalyst will make it possible to operate under milder process conditions (energy saving) or to increase the lifespan of a catalyst between regenerations (cycle length).

10 WO 96/41848 describes a process wherein a hydrotreating catalyst comprising a Group VIII hydrogenation metal oxide and a Group VI hydrogenation metal oxide on a carrier is activated by contacting it with an additive which is at least one compound selected from the group of compounds comprising at least two
15 hydroxyl groups and 2-10 carbon atoms, and the (poly)ethers of these compounds, after which the catalyst is dried under such conditions that the additive substantially remains in the catalyst. The additives described in this reference include sugars and various ethers and polyethers. The catalyst may be either a fresh (not used) catalyst or a used catalyst which has been regenerated.

20

WO 01/02092 describes a process for regenerating and activating a used
additive-based catalyst by contacting it with an oxygen-containing gas at a maximum temperature of 500°C, followed by activating it by contacting with an organic additive, if necessary followed by drying at such a temperature that at
25 least 50% of the additive is maintained in the catalyst. The preferred additives described in this reference are those selected from the group of compounds comprising at least two oxygen-containing moieties and 2-10 carbon atoms and the compounds built up from these compounds. Cited examples include acids, aliphatic-dialcohols, ethers thereof, sugars, and N-containing compounds.

30

It has now been found, however, that the process described in these references can be improved if the catalyst is activated with the combination of an acid and

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a specified additive. Accordingly, the present invention pertains to a process for activating an hydrotreating catalyst comprising a Group VIB metal oxide and a Group VIII metal oxide which process comprises containing the catalyst with an acid and an organic additive which has a boiling point in the range of 80-500°C and a solubility in water of at least 5 grams per liter (20°C, atmospheric pressure), optionally followed by drying under such conditions that at least 50% of the acid and additive is maintained in the catalyst. The process of the present invention is suitable for activating fresh hydrotreating catalysts, but also for activating catalysts which have been used in the hydrotreating of hydrocarbon feeds.

The present invention also relates to a hydrotreating catalyst comprising a Group VIII metal oxide and a Group VI metal oxide, which catalyst additionally comprises an acid and an organic additive which has a boiling point in the range of 80-500°C and a solubility in water of at least 5 grams per liter (20°C, atmospheric pressure). This catalyst can be obtained by the process of the present invention.

The starting material for the process according to the invention is a hydrotreating catalyst comprising a Group VIII hydrogenation metal oxide and a Group VI hydrogenation metal oxide, further indicated as an oxidic catalyst. Generally at least 80% of all of the Group VIII and Group VI hydrogenation metal present in the catalyst is in the oxidic form, preferably at least 90%, more preferably at least 95%, still more preferably at least 98%. The percentage of metals present in the catalyst in oxidic form can be determined by X-ray fluorescence spectroscopy (XRF) or inductively coupled plasma spectroscopy (ICP).

The oxidic hydrotreating catalyst used as starting material in the process according to the invention may thus be an oxidic hydrotreating catalyst prepared by a process in which hydrogenation metal components are composited with a carrier, after which the composite material is subjected to a calcination step to convert the hydrogenation metal components into their oxides. However, it may

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also be a used hydrotreating catalyst which has been regenerated by removing coke therefrom. In this case the catalyst may or may not have contained an additive before its first use.

- 5 The starting catalyst comprises a Group VIII metal oxide and a Group VI metal oxide, generally on a carrier. As Group VI metals may be mentioned molybdenum, tungsten, and chromium, with molybdenum or tungsten being preferred. Molybdenum is especially preferred. Group VIII metals include nickel, cobalt, and iron. Nickel, cobalt, or their combination is preferred. The catalyst
10 usually has a metal content in the range of 0,1 to 50 wt.%, calculated as oxides on the overall weight of the catalyst. The Group VI metal component is generally present in an amount of 5-40 wt.%, calculated as trioxide, preferably 10-35 wt.%, more preferably 15-30 wt.%. The Group VIII metal component is generally present in an amount of 1-10 wt.%, preferably 2-8 wt.%, calculated as
15 oxide. If so desired, the catalyst may also contain other components, such as phosphorus, halogens, and boron. Particularly, the presence of phosphorus in an amount of 1-10 wt.%, calculated as P_2O_5 , to improve the hydrodenitrogenation activity of the catalyst may be preferred.

The catalyst carrier may comprise the conventional oxides, e.g., alumina, silica,
20 silica-alumina, alumina with silica-alumina dispersed therein, silica-coated alumina, magnesia, zirconia, boria, and titania, as well as mixtures of these oxides. As a rule, preference is given to the carrier being of alumina, silica-alumina, alumina with silica-alumina dispersed therein, or silica-coated alumina. Special preference is given to alumina and alumina containing up to 10 wt.% of
25 silica. A carrier containing a transition alumina, for example an eta, theta, or gamma alumina is preferred within this group, wherein a gamma-alumina carrier is most especially preferred.

The catalyst's pore volume (measured via mercury penetration, contact angle 140°, surface tension of 480 dyn/cm) is not critical to the process according to
30 the invention and will generally be in the range of 0,2 to 2 ml/g, preferably 0.4-1 ml/g. The specific surface area is not critical to the process according to the invention either and will generally be in the range of 50 to 400 m²/g (measured

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using the BET method). Preferably, the catalyst will have a median pore diameter in the range of 7-15 nm, as determined by mercury porosimetry, and at least 60% of the total pore volume will be in the range of ± 2 nm from the median pore diameter.

- 5 The catalyst is employed in the conventional manner in the form of spheres or extrudates. Examples of suitable types of extrudates have been disclosed in the literature (see, *int. al.*, US 4 028 227). Highly suitable for use are cylindrical particles (which may be hollow or not) as well as symmetrical and asymmetrical polylobed particles (2, 3 or 4 lobes).

10

In the process according to the invention the oxidic hydrotreating catalyst is contacted with an acid and an organic additive having a boiling point in the range of 80-500°C and a solubility in water of at least 5 grams per liter.

- 15 The acid and the additive are incorporated into the catalyst in the liquid form by impregnation. For the acid this will generally mean it being in the dissolved state. For the additive, whether or not a solvent is necessary will depend on its properties. If the additive is sufficiently fluid to enter the catalyst's pores without a solvent being present, a solvent can be dispensed with. Generally, however, a solvent will be used. The solvent generally is water, although other compounds, 20 such as methanol, ethanol, and other alcohols may also be suitable, depending on the nature of the additive and the acid.

- In a preferred embodiment of the process of the invention, the catalyst is subjected to an aging step after the incorporation of the acid. The aging step is 25 effected while the catalyst is still wet, that is, before the solvent is removed from the catalyst. It has been found that the aging step is particularly advantageous when a catalyst which has been used in the hydrotreating of hydrocarbon feed is to be reactivated. The aging step generally takes at least 15 minutes. After aging more than 48 hours, no additional activity improvement is observed. The 30 aging time applied in the aging step is a function of temperature. Generally, the aging time decreases with increasing aging temperature. If the aging step is carried out at a temperature between 0° and 50°C, the aging time is generally at

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least 1 hour, preferably at least 2 hours, more preferably at least 6 hours. If the aging step is carried out at a temperature above 50°C, the aging time is generally at least 0.5 hours, preferably at least 1 hour, more preferably at least 2 hours. It is also possible to effect the aging step at a temperature of above
5 100°C under hydrothermal conditions for a period of over 15 minutes. The aging step can also be performed by heating the catalyst with microwaves or induction heating.

If solvent has been used to incorporate the additive and/or the acid into the
10 catalyst, the catalyst may be dried after the impregnation step has been completed to remove at least part of the solvent, generally at least 50%, preferably at least 70%, more preferably at least 80%. It is essential to the process according to the invention that any drying step is effected in such a manner that at least part of the additive remains in the catalyst. In
15 consequence, the drying conditions to be applied depend heavily on the temperature at which the specific additive boils or decomposes. In the context of the present invention, the drying step should be carried out under such conditions that at least 50%, preferably 70 %, more preferably 90% of the additive incorporated into the catalyst in the impregnation step is still present in
20 the catalyst after the drying step. Of course, it is preferred to keep as much additive as possible in the catalyst during the drying step, but with the more volatile compounds evaporation during the drying step cannot always be avoided. The drying step may, e.g., be carried out in air, under vacuum, or in inert gas. Generally, it is advantageous to have a drying temperature below
25 220°C, although a higher or lower temperature may be necessary, depending on the nature of the additive.

The acid and the additive may be incorporated into the catalyst simultaneously or sequentially in no particular order.

30 In one embodiment, the starting material is contacted with a solution of an acid, optionally followed by drying. Then, the organic additive is incorporated into the catalyst, optionally followed by a drying step under such conditions that at least

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50% of the additive remains in the catalyst. If in this embodiment an aging step is carried out, it may be performed before or after the organic additive has been incorporated into the catalyst composition.

In another embodiment, in a first step the organic additive is incorporated into the starting material, optionally followed by a drying step under such conditions that at least 50% of the additive remains in the catalyst. Then, the resulting material is contacted with a solution of an acid, optionally followed an aging step and/or by drying under such conditions that at least 50% of the additive remain in the catalyst.

The advantage of incorporating the acid and the additive into the catalyst in separate steps is that the properties of the impregnation solutions may be tailored to meet the requirements of the acid and the additive. Nevertheless, for reasons of efficiency, it is preferred to contact the starting catalyst with a single impregnation solution comprising both the acid and the additive, optionally followed by an aging step and/or a drying step under such conditions that at least 50% of the additive remains in the catalyst.

The acid may be an inorganic or an organic acid. Inorganic acids are defined as acidic components which do not contain a carbon atom. Examples of inorganic acids include HCl, H₂SO₄, HNO₃, H₃PO₄, (NH₄)H₂PO₄, (NH₄)₂HPO₄, H₂PHO₃, H₂P₂H₂O₅, and H_(n+2)P_nO_(3n+1). Within the group of inorganic acids, the phosphorus-containing acids are preferred, since phosphorus in itself can positively influence the activity of a hydrotreating catalyst. H₃PO₄ is particularly preferred.

One disadvantage to the use of inorganic acids is that the counterion, e.g., chloride, sulfate, or nitrate, is incorporated into the catalyst composition. If the activation process of the present invention is repeated a number of times this may lead to an undesirable build-up of these counterions in the catalyst composition. However, depending on the nature of the counterion and on the number of regeneration steps envisaged, this may not be a problem.

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In the context of the present specification, an organic acid is defined as a compound comprising at least one carboxylic group (COOH). The organic acid is generally a carboxylic acid comprising at least one carboxylgroup and 1-20 carbon atoms (carbon atoms in the carboxylgroup included). Suitable acids include acetic acid, citric acid, malic acid, maleic acid, formic acid, glycolic acid, hydroxypropionic acid, hydroxybutyric acid, hydroxyhexanoic acid, tartaric acid, glyceric acid, gluconic acid, oxalic acid, malonic acid, polyacrylic acid, ascorbic acid, and N-containing acids like EDTA and CyDTA (1,2-cyclohexanediaminetetra-acetic acid) etc. Within this group, compounds comprising at least two carboxylic groups are preferred. For combined costs and activity, citric acid, malic acid, maleic acid, malonic acid, and tartaric acid are preferred. Citric acid is particularly preferred.

The organic additive used in combination with the acid in the process according to the invention is an organic compound, viz. a compound comprising at least one carbon atom and at least one hydrogen atom, with a boiling point in the range of 80-500°C and a solubility in water of at least 5 grams per liter at room temperature (20°C) (atmospheric pressure). Preferably the additive is an oxygen- or nitrogen-containing compound.

The boiling point of the additive is preferably in the range of 100-400°C, more preferably 150-350°C. The boiling point of the additive is balanced between on the one hand the desire that the additive remain on the catalyst during the preparation process, including the drying step, and on the other hand the need for the additive to be removed from the catalyst during catalyst use or sulfidation.

The solubility of the additive is at least 5 grams per liter at room temperature, preferably at least 10 grams. It is noted that the solubility requirement for the additive has a two-fold basis. In the first place, compounds meeting this solubility requirement are convenient to apply in impregnation solutions. Additionally it has appeared that compounds which meet these solubility requirements somehow interact with the metal components present in the catalyst leading to an increase in activity of the final product.

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Within this definition, various groups of additives may be distinguished.

A first, at this point in time preferred, group of additives is the group of organic compounds comprising at least two oxygen atoms and 2-20 carbon atoms, preferably 2-10 carbon atoms and the compounds built up from these compounds. Organic compounds selected from the group of compounds comprising at least two oxygen-containing moieties, such as a carboxyl, carbonyl or hydroxyl moieties, and 2-10 carbon atoms and the compounds built up from these compounds are preferred. Examples of suitable compounds include butanediol, pyruvic aldehyde, glycolic aldehyde, and acetaldehyde. At this point in time, preference is given within this group to an additive that is selected from the group of compounds comprising at least two hydroxyl groups and 2-10 carbon atoms per molecule, and the (poly)ethers of these compounds. Suitable compounds from this group include aliphatic alcohols such as ethylene glycol, propylene glycol, glycerin, trimethylol ethane, trimethylol propane, etc. Ethers of these compounds include diethylene glycol, dipropylene glycol, trimethylene glycol, triethylene glycol, tributylene glycol, tetraethylene glycol, tetrapentylene glycol. This range can be extrapolated to include polyethers like polyethylene glycol. Other ethers which are suitable for use in the present invention include ethylene glycol monobutyl ether, diethylene glycol monomethyl ether, diethylene glycol monoethyl ether, diethylene glycol monopropyl ether, and diethylene glycol monobutyl ether. Of these, ethylene glycol, diethylene glycol, triethylene glycol, tetraethylene glycol, propylene glycol, dipropylene glycol, and polyethylene glycol with a molecular weight between 200 and 600 are preferred. Another group of compounds comprising at least two hydroxyl groups and 2-10 carbon atoms per molecule are the saccharides. Preferred saccharides include monosaccharides such as glucose and fructose. Ethers thereof include disaccharides such as lactose, maltose, and saccharose. Polyethers of these compounds include the polysaccharides. The organic compounds of this group are preferably substantially saturated, as is evidenced by a iodine number of less than 60, preferably less than 20.

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A second group of organic additives suitable for use in the present invention are those compounds comprising at least one covalently bonded nitrogen atom and at least one carbonyl moiety. This type of organic compound preferably
5 comprises at least two carbonyl moieties. It is preferred that at least one carbonyl moiety is present in a carboxyl group. It is furthermore preferred that at least one nitrogen atom is covalently bonded to at least two carbon atoms. A preferred organic compound satisfies formula (I) or (II)

10 (R₁R₂)N - R₃ - N(R₁'R₂') (I)

N(R₁R₂R₁') (II)

wherein R₁, R₂, R₁' and R₂' are independently selected from alkyl, alkenyl, and
15 allyl, with up to 10 carbon atoms optionally substituted with one or more groups selected from carbonyl, carboxyl, ester, ether, amino, or amido. R₃ is an alkylene group with up to 10 carbon atoms which may be interrupted by -O- or -NR₄-. R₄ is selected from the same group as indicated above for R₁. The R₃ alkylene group may be substituted with one or more groups selected from
20 carbonyl, carboxyl, ester, ether, amino, or amido. As has been set out above, it is essential that the organic compound of formula (I) or (II) comprises at least one carbonyl moiety.

Preferably, at least two of R₁, R₂, R₁' and R₂' (formula (I)) and at least two of R₁, R₂ and R₁' (formula (II)) have the formula - R₅ - COOX, wherein R₅ is an
25 alkylene group having 1-4 carbon atoms, and X is hydrogen or another cation, such as an ammonium, sodium, potassium and/or lithium cation. If X is a multivalent cation, one X can adhere to two or more - R₅ - COO groups. Typical examples of a compound of formula (I) are ethylene diamine(tetra)acetic acid (EDTA), hydroxyethylene diamine triacetic acid, and diethylene triamine
30 pentaacetic acid. A typical example of a compound of formula (II) is nitrilotriacetic acid (NTA). For reasons of solubility, the salts of these compounds may be preferred.

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It is noted that the above-mentioned description of the organic additive encompasses various acidic components. Where in the context of the present invention the combination of an acid and an organic additive is discussed, this
5 means that (at least) two different compounds are used wherein one is an acid and wherein the other meets the requirements as to solubility and boiling point placed on the organic additive. The latter compound may or may not also be acidic.

10 From an environmental point of view, it is preferred to employ additives which are essentially free of sulfur. Further, sulfur-containing additives generally are not stable in relation to oxygen. Therefore, if sulfur-containing additives were to be employed, all subsequent process steps would have to be carried out under an inert atmosphere. Also for this reason it is preferred to employ sulfur-free
15 additives. This goes both for the acid and for the organic additive.

A single compound as well as a combination of compounds may be used as additive.

20 The total amount of acid and additive used in the process according to the invention, and the total amount of acid and additive present in the catalyst according to the invention is at least 0.01, preferably at least 0.05, more preferably at least 0.1 mole of total of acid and additive per mole of the total of Group VIB and Group VIII metals. Generally, the molar ratio will be at most 3,
25 preferably at most 2.

If an inorganic acid is used, the amount of acid is generally between 0.01 and 1 mole per mole of total of Group VIB and Group VIII metals, preferably between 0.05 and 0.5 mole mole per mole of total of Group VIB and Group VIII metals.

If an organic acid is used, the amount of acid is generally between 0.01 and 1
30 mole per mole of total of Group VIB and Group VIII metals, preferably between 0.05 and 0.5 mole per mole of total of Group VIB and Group VIII metals.

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The amount of organic additive is generally between 0.1 and 2,5 mole per mole of total of Group VIB and Group VIII metals, preferably between 0.15 and 1 mole per mole of total of Group VIB and Group VIII metals, more preferably between 0.2 and 1 mole per mole of total of Group VIB and Group VIII metals.

- 5 The molar ratio between acid and additive is generally 0.01-10:1, preferably 0.1-5:1, more preferably 0.15-3:1.

10 In general, if the total amount of acid and additive added is too low, the advantageous effect of the present invention will not be obtained. On the other hand, the addition of an exceptionally large total amount of acid and additive will not improve the effect of the present invention. As will be clear to the person skilled in the art, the exact amount of acid and additive to be used in a specific situation will depend upon a variety of parameters including the metals content of the catalyst, the pore volume and pore size distribution of the catalyst, the
15 nature of the acid and additive, the solvent to be used in the impregnation solution, the impregnation conditions, etc. It is well within the scope of the person skilled in the art to determine the optimum amount of acid and additive to be used in each specific situation, taking the above-mentioned variables into account.

20

The present invention also pertains to the hydrotreating catalyst which comprises a Group VIII metal oxide and a Group VI metal oxide on a carrier, which catalyst additionally comprises an acid and an organic additive as described above.

25

The dried additive-containing hydrotreating catalyst of the present invention may be subjected to a sulfiding step before it is used in the hydrotreating of hydrocarbon feeds, but, as has been explained before, this is not necessary. If it is decided to sulfide the catalyst before use, this can be done in one of the ways
30 known in the art. For example, it is possible to contact the catalyst with inorganic or organic sulfur compounds, such as hydrogen sulfide, elemental sulfur, or organic polysulfides, or to sulfide the catalyst by contacting it with a

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hydrocarbon feed to which a sulfur compound has been added. All of this will be known to the skilled person as catalyst sulfiding or presulfiding.

The present catalysts can be used in the hydrotreating of a wide range of feeds. To effect one or more of hydrodesulfurisation, hydrodenitrogenation, and
5 hydrodearomatisation. Examples of suitable feeds include middle distillates, kero, naphtha, vacuum gas oils, and heavy gas oils. The catalyst is particularly suitable for use in ultra-deep hydrodesulfurisation, viz. hydrodesulfurisation to a product sulphur content below 200 ppm, more in particular to product sulfur
10 content below 50 ppm. The conventional process conditions, such as temperatures in the range of 250°-450°C, pressures in the range of 5-250 bar, space velocities in the range of 0,1-10 h⁻¹, and H₂/oil ratios in the range of 50-2000 Nl/l, can be applied here.

As indicated above, the catalyst to be activated in the process according to the
15 invention is either a fresh hydrotreating catalyst or used and regenerated hydrotreating catalyst.

The fresh oxidic hydrotreating catalyst suitable for use as starting material in the process according to the invention are known in the art. They may be obtained, e.g., as follows. A carrier precursor is prepared, e.g., in the case of alumina, in
20 the form of an alumina hydrogel (boehmite). After it has been dried or not, e.g., by means of spray-drying, it is shaped into particles, for example by extrusion. The shaped particles are calcined at a temperature in the range of 400° to 850°C, resulting, in the case of alumina, in a carrier containing a transition alumina, e.g., a gamma, theta, or eta-alumina. Then, suitable amounts of
25 precursors for the hydrogenation metals and the optional other components, such as phosphorus, are deposited on the catalyst, for example in the form of an aqueous solution. In the case of Group VI metals and Group VIII metals, the precursors may be ammonium molybdate, ammonium tungstenate, cobalt nitrate and/or nickel nitrate. Suitable phosphorus component precursors include
30 phosphoric acid and the various ammonium hydrogen phosphates. After an optional drying step at a temperature in the range of 25°-200°C, the resulting material is calcined at a temperature in the range of 350°-750°C, more in

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particular 425 to 600°C, to convert all metal component precursors, and the optional other component precursors to form oxide components.

It will be clear to the skilled man that there can be a wide number of variations on this method. Thus, it is possible to apply a plurality of impregnating steps, the impregnating solutions to be used containing one or more of the component precursors that are to be deposited, or a portion thereof. Instead of impregnating techniques there can be used dipping methods, spraying methods, etc. With multiple impregnation, dipping, etc. drying and/or calcining may be carried out in between. Alternatively, one or more component precursors can be mixed wholly or in part with the carrier prior to the shaping step being carried out. In such an embodiment the component precursor material, or a portion thereof, can be deposited on the not yet shaped carrier material, say, a powder, by means of, e.g., impregnation, after which the shaping step is carried out. Alternatively, it is possible to intimately mix one or more component precursors wholly or in part with the carrier material during the shaping step rather than prior to it. Techniques that are suitable for this process are co-pelletisation and co-extrusion. It is recommended that the group VIB metal component precursor employed in this process be molybdenum trioxide. Oxidic hydrotreating catalysts and the processes for preparing them are part of the common general knowledge of the skilled person and described in, e.g., US 4 738 767, US 4 062, 809, US 4 500 424, GB 1 504 586, US 4 212 729, US 4 326 995, US 4 051 021, US 4 066 574, EP-A 0 469 675.

The activation process of the present invention is also applicable to the catalyst which has been used in the hydrotreating of hydrocarbon feeds and subsequently regenerated.

The present invention also pertains to a combined process for regenerating and activating a used hydrotreating catalyst wherein a used hydrotreating catalyst comprising a Group VIB metal oxide and a Group VIII metal oxide is first subjected to a regeneration step to remove carbonaceous and sulfurous deposits, followed by activating the thus obtained material by contacting it with an acid and an organic additive which has a boiling point in the range of 80-

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500°C and a solubility in water of at least 5 grams per liter (20°C, atmospheric pressure).

The regeneration step of the process according to the invention is carried out by
5 contacting the used additive-based catalyst with an oxygen-containing gas
under such conditions that after regeneration, the carbon content of the catalyst
generally is below 3 wt.%, preferably below 2 wt.%, more preferably below 1
wt.%. After regeneration, the sulfur content of the catalyst generally is below 2
wt.%, preferably below 1 wt.%. Before the regeneration step, the carbon content
10 of the catalyst generally is above 5 wt.%, typically between 5 and 25 wt.%. The
sulfur content of the catalyst before the regeneration step generally is above 5
wt.%, typically between 5 and 20 wt.%.

The maximum catalyst temperature during the regeneration step will be
15 governed by the properties of the catalyst to be regenerated and by process
constraints, a higher maximum temperature being preferred in principle
because this makes it possible to reduce the regeneration time. However, a
high regeneration temperature carries the risk of damaging the catalyst.
Catalysts with a higher metal content will generally require a lower maximum
20 catalyst temperature than catalysts with a lower metal content. Generally, the
maximum catalyst temperature during the regeneration process is at most
650°C, preferably at most 575°C, more preferably at most 550°C, still more
preferably at most 525°C.

The maximum catalyst temperature during the regeneration process generally is
25 at least 300°C, preferably at least 350°C, more preferably at least 400°C, still
more preferably at least 450°C.

It is noted that in this specification any temperature given relates to the
temperature of the catalyst, except when explicitly indicated otherwise. The
30 catalyst temperature can be determined in any way known to the skilled person,
e.g., by way of appropriately placed thermocouples.

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It is preferred for the regeneration step in the presence of oxygen to be carried out in two steps, namely a first lower-temperature step and a second higher-temperature step. In the first, lower-temperature step, the catalyst is contacted with an oxygen-containing gas at a temperature of 100 to 370°C, preferably 175 to 370°C. In the second, higher-temperature regeneration step, the catalyst is contacted with an oxygen-containing gas at a temperature of In the second, higher-temperature regeneration step, the catalyst is contacted with an oxygen-containing gas at a temperature of 300 to 650°C, preferably 320 to 550°C, still more preferably 350-525°C. The temperature during the second step is higher than the temperature of the first step discussed above, preferably by at least 10°C, more preferably by at least 20°C. The determination of appropriate temperature ranges is well within the scope of the skilled person, taking the above indications into account.

It is preferred for the catalyst to be regenerated in a moving bed process, preferably, if applicable, at a bed thickness of 1-15 cm. In the context of the present specification, the term "moving bed" is intended to refer to all processes wherein the catalyst is in movement as compared to the unit, including ebullated bed processes, fluidised processes, processes in which the catalyst is rotated through a unit, and all other processes wherein the catalyst is in movement.

The duration of the regeneration process including stripping will depend on the properties of the catalyst and the exact way in which the process is carried out, but will generally be between 0.25 and 24 hours, preferably between 2 and 16 hours.

The regenerated catalyst will be contacted with the acid and additive in the process according to the invention as has been described above.

The invention will be illustrated by the following examples, but is not intended to be limited thereto or thereby.

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Example 1: Activating used catalyst with citric acid and polyethylene glycol

5 A hydroprocessing catalyst that had been used for 12 months on the hydrotreating of a LGO feedstock was regenerated as follows. The catalyst contained 23.2 wt.% of carbon, 9 wt.% of sulfur, and 22.4 wt.% of hydrocarbon. In a first step, the hydrocarbon was removed from the catalyst by stripping. Then, the catalyst was regenerated by contacting it with air at a temperature of 490-500°C for a period of 24 hours. The regenerated starting material thus
10 obtained contained molybdenum, cobalt, and phosphorus on an alumina carrier.

The regenerated catalyst was impregnated via pore volume impregnation with an impregnation solution containing citric acid and polyethylene glycol in an amount of 0.1 mole acid per mole of hydrogenation metals and 0.5 mole
15 polyethylene glycol per mole of hydrogenation metals, respectively, and subsequently dried overnight at a temperature of 120°C.

The catalyst was tested on a LGO feedstock having the following properties:

| | |
|-----------------------------------|--------|
| S (wt.%) | 1.4837 |
| N (wt. ppm) | 170 |
| density (15°C, g/ml) | 0.8609 |
| viscosity (50°C, cSt) | 4.11 |
| H (wt.%) | 12.90 |
| Distillation Data (ASTM D 86, °C) | |
| Initial boiling point | 182 |
| 10 vol.% | 288 |
| 50 vol.% | 325 |
| 90 vol.% | 364 |
| Final boiling point | 375 |

20 The catalyst was tested at a liquid hourly space velocity (LHSV) of 1.5 h⁻¹ and a hydrogen to oil ratio of 200 NI/I at various conditions with varying temperature and pressure. Each condition was maintained for two days. The catalyst was

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tested side by side with a fresh catalyst. The test conditions and activities of the two catalysts are given below.

| | T (°C) | PpH2 (MPa) | fresh catalyst RVA-HDS (ppm S) | reactivated catalyst RVA-HDS (ppm S) |
|---|--------|------------|-----------------------------------|---|
| 1 | 345 | 5.1 | 100 (42) | 93 (50) |
| 2 | 350 | 5.1 | 100 (28) | 93 (33) |
| 3 | 350 | 3.0 | 100 (159) | 98 (165) |
| 4 | 350 | 5.1 | 100 (55) | 103 (52) |

- 5 As can be seen from the above table, the activity of the spent and reactivated catalyst had been brought back to an activity which is substantially as high as the activity of the fresh catalyst. In comparison, the activity of the spent catalyst after regeneration was 70% as compared with its initial activity.

10 Example 2: Activating used catalyst with phosphoric acid and polyethylene glycol

A hydroprocessing catalyst that had been used for 24 months on the hydrotreating of a LGO feedstock was regenerated as follows. The catalyst
15 contained 25.4 wt.% of carbon, 8.8 wt.% of sulfur, and 18.3 wt.% of hydrocarbon. In a first step, the hydrocarbon was removed from the catalyst by stripping. Then, the catalyst was regenerated by contacting it with air at a temperature of 450°C for a period of 24 hours. The regenerated starting material thus obtained contained molybdenum, cobalt, and phosphorus on an
20 alumina carrier.

The regenerated catalyst was impregnated via pore volume impregnation with an impregnation solution containing phosphoric acid and polyethylene glycol in an amount of 0.2 mole polyethylene glycol per mole of hydrogenation metals,
25 and subsequently dried overnight at a temperature of 120°C. The final catalyst contained 4.8 wt.% of phosphorus, calculated as P₂O₅ while the original catalyst contained 1.6 wt.%.

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The catalyst was tested on a LGO feedstock having the following properties:

| | |
|-----------------------------------|--------|
| S (wt.%) | 1.323 |
| N (wt. ppm) | 110 |
| density (15°C, g/ml) | 0.8574 |
| Distillation Data (ASTM D 86, °C) | |
| Initial boiling point | 183 |
| 10 vol.% | 253 |
| 50 vol.% | 298 |
| 90 vol.% | 360 |
| Final boiling point | 380 |

The catalyst was tested at a hydrogen partial pressure of 3.92 MPa, a liquid hourly space velocity (LHSV) of 1.5 h⁻¹, and a hydrogen to oil ratio of 200 NI/l at various conditions with varying temperatures. Each condition was maintained for two days. The catalyst was tested side by side with a fresh catalyst. The test conditions and activities of the two catalysts are given below.

| | T (°C) | fresh catalyst RVA-HDS (ppm S) | reactivated catalyst RVA-HDS (ppm S) |
|---|--------|-----------------------------------|---|
| 1 | 340 | 100 (114) | 98 (120) |
| 2 | 350 | 100 (39) | 99 (41) |
| 3 | 330 | 100 (406) | 101 (398) |

10

As can be seen from the above table, the activity of the spent and reactivated catalyst had been brought back to an activity which is substantially as high as the activity of the fresh catalyst. In comparison, the relative volume activity for HDS of the spent catalyst after having been used for 24 months in the hydrotreating of a hydrocarbon oil had decreased to 53% as compared with its initial relative volume activity.

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Example 3: Activating used catalyst with citric acid and polyethylene glycol

A hydroprocessing catalyst that had been used in the hydrotreating of a hydrocarbon feedstock was regenerated as follows. The catalyst contained 17.9

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wt.% of carbon and 10.7 wt.% of sulfur. The catalyst was regenerated by contacting it with air at a temperature of 500-530°C. The regenerated starting material thus obtained contained molybdenum, nickel, and phosphorus on an alumina carrier.

5

The regenerated catalyst was impregnated via pore volume impregnation with an impregnation solution containing citric acid and polyethylene glycol in an amount of 0.09 mole acid per mole of hydrogenation metals and 0.22 mole polyethylene glycol per mole of hydrogenation metals, respectively, and subsequently aged for 24 hours while wet and dried overnight at a temperature of 90°C.

10

The catalyst was tested on a CGO feedstock having the following properties:

| | |
|-------------------------------------|--------|
| S (wt.%) | 2.44 |
| N (wt. ppm) | 3546 |
| density (15°C, g/ml) | 0.9173 |
| H (wt.%) | 11.7 |
| Distillation Data (ASTM D 1160, °C) | |
| Initial boiling point | 176°C |
| 10 vol.% | 250°C |
| 50 vol.% | 332°C |
| 90 vol.% | 412°C |
| Final boiling point | 503°C |

15 The catalyst was tested at a temperature of 387°C, a pressure of 100 bar, a liquid hourly space velocity (LHSV) of 2.1 h⁻¹, and a hydrogen to oil ratio of 1000 Nl/l. The hydrodenitrogenation activity of the regenerated and reactivated catalyst was 90% of the hydrodenitrogenation activity of the fresh catalyst. In comparison, the activity of the spent catalyst after regeneration but before
20 reactivation was about 60% of the activity of the fresh catalyst.

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CLAIMS

1. A process for activating an hydrotreating catalyst comprising a Group VIB metal oxide and a Group VIII metal oxide which process comprises containing the catalyst with an acid and an organic additive which has a boiling point in the range of 80-500°C and a solubility in water of at least 5 grams per liter (20°C, atmospheric pressure), optionally followed by drying under such conditions that at least 50% of the additive is maintained in the catalyst.
2. A process according to claim 1, wherein the hydrotreating catalyst is a fresh hydrotreating catalyst.
3. A process according to claim 1 wherein the hydrotreating catalyst is a used hydrotreating catalyst which has been regenerated.
4. The process of claim 3 wherein the catalyst composition containing the acid is subjected to an aging step while wet.
5. A process according to any one of the preceding claims wherein the acid is an inorganic acid, preferably a phosphorus-containing inorganic acid.
6. A process according to any one of the preceding claims 1-4 wherein the acid is a carboxylic acid comprising at least one carboxylgroup and 1-20 carbon atoms.
7. A process according to claim 6 wherein the acid is citric acid.
8. A process according to any of the preceding claims wherein the additive is an organic oxygen- or nitrogen- containing compound, with a boiling

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point in the range of 100-400°C and a solubility in water of at least 5 grams per liter at room temperature (20°C) (atmospheric pressure).

- 5 9. A process according to claim 8 wherein the additive is selected from the group of compounds comprising at least two hydroxyl groups and 2-10 carbon atoms per molecule, and the (poly)ethers of these compounds.
- 10 10. A hydrotreating catalyst comprising a Group VIII metal oxide and a Group VI metal oxide, which catalyst additionally comprises an acid and an organic additive which has a boiling point in the range of 80-500°C and a solubility in water of at least 5 grams per liter (20°C, atmospheric pressure).
- 15 11. A process for hydrotreating a hydrocarbon feed in which a hydrocarbon feed is contacted under hydrotreating conditions with a catalyst according to claim 10, which optionally has been (pre)sulfided before it is contacted with the hydrocarbon feed.

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ABSTRACT

The invention pertains to a process for activating an hydrotreating catalyst
5 comprising a Group VIB metal oxide and a Group VIII metal oxide which
process comprises containing the catalyst with an acid and an organic additive
which has a boiling point in the range of 80-500°C and a solubility in water of at
least 5 grams per liter (20°C, atmospheric pressure), optionally followed by
drying under such conditions that at least 50% of the additive is maintained in
10 the catalyst. The hydrotreating catalyst may be a fresh hydrotreating catalyst or
a used hydrotreating catalyst which has been regenerated.

The use of the combination of an acid and an additive results in a catalyst with
higher activity as compared to the use of an acid alone or an additive alone.

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